FDA 510(k) Premarket Notification

K103176 FREQUENCER

CLEARED

Jan 26, 2011
90 days to decision

K103176 · Product code: **BYI** · Anesthesiology
Source: https://www.510kdatabase.net/k103176/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Percussor, Powered-electric (BYI)
Date received	Oct 28, 2010
Decision date	Jan 26, 2011
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Dymedso, Inc.**
Location	Springfield, VT, US
Contact	Jean Bigoney
510(k) history	3 submissions · 3 cleared · 2007-2011